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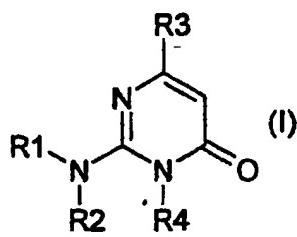
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(54) Title: 2-AMINO-3-(ALKYL)-PYRIMIDONE DERIVATIVES AS GSK3 β .INHIBITORS



(57) Abstract: A pyrimidone derivative represented by formula (I) or a salt thereof, wherein: R1 represents a hydrogen atom or a C₁₋₆ alkyl group which may be substituted by a C₆₋₁₀ aryl group; R2 represents a C₁₋₁₀ alkyl group which may be substituted, a C₂₋₆ alkenyl group which may be substituted, a C₃₋₆ alkynyl group which may be substituted, a C₃₋₆ cycloalkyl group which may be substituted, or a C₆₋₁₀ ARYL group which may be substituted; or R1 and R2 form together a C₂₋₆ alkylene group which may be substituted; or R1 and R2 form together a chain of formula -(CH₂)₂-X-(CH₂)₂- or -(CH₂)₂-X-(CH₂)₃- where X represents an oxygen atom, a sulfur atom, or a nitrogen atom which may be substituted; R3 represents a 2, 3 or 4-pyridyl group optionally substituted by a C₁₋₄ alkyl group, C₁₋₄ alkoxy group or halogen atom; and R4 represents a C₁₋₁₀ alkyl group optionally substituted by a hydroxyl group, amino, C₁₋₆ monoalkylamino group, C₂₋₁₂ dialkylamino group or C_{7-10?} aryl group which may be substituted. And a medicament comprising the said derivative or a salt thereof as an active ingredient which is used for preventive and/or therapeutic treatment of a neurodegenerative disease caused by abnormal activity of GSK3 β such as Alzheimer's disease.

substituted by a hydroxyl group, amino, C₁₋₆ monoalkylamino group, C₂₋₁₂ dialkylamino group or C_{7-10?} aryl group which may be substituted. And a medicament comprising the said derivative or a salt thereof as an active ingredient which is used for preventive and/or therapeutic treatment of a neurodegenerative disease caused by abnormal activity of GSK3 β such as Alzheimer's disease.

SPECIFICATION

2-AMINO-3- (ALKYL) -PYRIMIDONE DERIVATIVES AS GSK3 β . INHIBITORS

5 Technical Field

The present invention relates to compounds that are useful as an active ingredient of a medicament for preventive and/or therapeutic treatment of neurodegenerative diseases caused by abnormal activity of GSK3 β .

10 Background Art

GSK3 β (glycogen synthase kinase 3 β) is a proline directed serine, threonine kinase that plays an important role in the control of metabolism, differentiation and survival. It was initially identified as an enzyme able to phosphorylate and hence inhibit glycogen synthase. It was later recognized that GSK3 β was identical to tau protein kinase 1 (TPK1), an enzyme that phosphorylates tau protein in epitopes that are also found to be hyperphosphorylated in Alzheimer's disease and in several taupathies.

Interestingly, protein kinase B (AKT) phosphorylation of GSK3 β results in a loss of its kinase activity, and it has been hypothesized that this inhibition may mediate some of the effects of neurotrophic factors. Moreover, phosphorylation by GSK3 β of β -catenin, a protein involved in cell survival, results in its degradation by an ubiquitination dependent proteasome pathway.

Thus, it appears that inhibition of GSK3 β activity may result in neurotrophic activity.

Indeed there is evidence that lithium, an uncompetitive inhibitor of GSK3 β , enhances neuritogenesis in some models and also increases neuronal survival, through the induction of survival factors such as Bcl-2 and the inhibition of the expression of proapoptotic factors such as P53 and Bax.

Recent studies have demonstrated that β -amyloid increases the GSK3 β activity and tau protein phosphorylation. Moreover, this hyperphosphorylation as well as the neurotoxic effects of β -amyloid are blocked by lithium chloride and by a GSK3 β antisense mRNA. These observations strongly suggest that GSK3 β may be the link between the two major pathological processes in Alzheimer's disease : abnormal APP (Amyloid Precursor Protein) processing and tau protein hyperphosphorylation.

Although tau hyperphosphorylation results in a destabilization of the neuronal cytoskeleton, the pathological consequences of abnormal GSK3 β activity are, most likely, not only due to a pathological phosphorylation of tau protein because,

as mentioned above, an excessive activity of this kinase may affect survival through the modulation of the expression of apoptotic and antiapoptotic factors. Moreover, it has been shown that β -amyloid-induced increase in GSK3 β activity results in the phosphorylation and, hence the inhibition of pyruvate dehydrogenase, a pivotal enzyme in energy production and acetylcholine synthesis.

Altogether these experimental observations indicate that GSK3 β may find application in the treatment of the neuropathological consequences and the cognitive and attention deficits associated with Alzheimer's disease, as well as other acute and chronic neurodegenerative diseases. These include, in a non-limiting manner, Parkinson's disease, tauopathies (e.g. frontotemporoparietal dementia, corticobasal degeneration, Pick's disease, progressive supranuclear palsy) and other dementia including vascular dementia; acute stroke and others traumatic injuries; cerebrovascular accidents (e.g. age related macular degeneration); brain and spinal cord trauma; peripheral neuropathies; retinopathies and glaucoma.

In addition GSK3 β may find application in the treatment of other diseases such as:
20 Non-insulin dependent diabetes (such as diabetes type II) and obesity; manic depressive illness; schizophrenia; alopecia; cancers such as breast cancer, non-small cell lung carcinoma, thyroid cancer, T or B-cell leukemia and several virus-induced tumors.

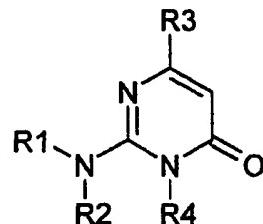
25 Disclosure of the Invention

An object of the present invention is to provide compounds useful as an active ingredient of a medicament for preventive and/or therapeutic treatment of neurodegenerative diseases. More specifically, the object is to provide novel 30 compounds useful as an active ingredient of a medicament that enables prevention and/or treatment of the diseases such as Alzheimer's.

Thus the inventors of the present invention have identified compounds possessing inhibitory activity against GSK3 β .

As a result, they found that compounds represented by the following 35 formula (I) had the desired activity and were useful as an active ingredient of a medicament for preventive and/or therapeutic treatment of the aforementioned diseases.

The present invention thus provides pyrimidone derivatives represented by formula (I) or salts thereof, solvates thereof or hydrates thereof:



(I)

5 Wherein :

R1 represents a hydrogen atom or a C₁₋₆ alkyl group which may be substituted by a C₆₋₁₀ aryl group;

R2 represents a C₁₋₁₀ alkyl group which may be substituted, a C₂₋₆ alkenyl group which may be substituted, a C₃₋₆ alkynyl group which may be substituted, a C₃₋₆

10 cycloalkyl group which may be substituted, or a C₆₋₁₀ ARYL group which may be substituted;

or R1 and R2 form together a C₂₋₆ alkylene group which may be substituted;

or R1 and R2 form together a chain of formula -(CH₂)₂-X-(CH₂)₂- or

-(CH₂)₂-X-(CH₂)₃- where X represents a oxygen atom, a sulfur atom, or a

15 nitrogen atom which may be substituted;

R3 represents a 2, 3 or 4-pyridyl group optionally substituted by a C₁₋₄ alkyl group, C₁₋₄ alkoxy group or halogen atom; and

R4 represents a C₁₋₁₀ alkyl group optionally substituted by a hydroxyl group, amino, C₁₋₆ monoalkylamino group, C₂₋₁₂ dialkylamino group or C₆₋₁₀ aryl group

20 which may be substituted.

According to another aspect of the present invention, there is provided a medicament comprising as an active ingredient a substance selected from the group consisting of the pyrimidone derivatives represented by formula (I) and the 25 physiologically acceptable salts thereof, and the solvates thereof and the hydrates thereof. As preferred embodiments of the medicament, there are provided the aforementioned medicament which is used for preventive and/or therapeutic treatment of diseases caused by abnormal GSK3 β activity, and the aforementioned medicament which is used for preventive and/or therapeutic treatment of neurodegenerative diseases and in addition other diseases such as: Non-insulin dependent diabetes (such as diabetes type II) and obesity; manic depressive illness; schizophrenia; alopecia; cancers such as breast cancer, non- 30

small cell lung carcinoma, thyroid cancer, T or B-cell leukemia and several virus-induced tumors.

As further preferred embodiments of the present invention, there are provided the
5 aforementioned medicament wherein the diseases are neurodegenerative
diseases and are selected from the group consisting of Alzheimer's disease,
Parkinson's disease, tauopathies (e.g. frontotemporoparietal dementia,
corticobasal degeneration, Pick's disease, progressive supranuclear palsy) and
other dementia including vascular dementia; acute stroke and others traumatic
10 injuries; cerebrovascular accidents (e.g. age related macular degeneration); brain
and spinal cord trauma; peripheral neuropathies; retinopathies and glaucoma, and
the aforementioned medicament in the form of pharmaceutical composition
containing the above substance as an active ingredient together with one or more
pharmaceutical additives.

15

The present invention further provides an inhibitor of GSK3 β activity comprising as an active ingredient a substance selected from the group consisting of the pyrimidone derivatives of formula (I) and the salts thereof, and the solvates thereof and the hydrates thereof.

20

According to further aspects of the present invention, there are provided a method for preventive and/or therapeutic treatment of neurodegenerative diseases caused by abnormal GSK3 β activity, which comprises the step of administering to a patient a preventively and/or therapeutically effective amount of a substance selected from the group consisting of the pyrimidone derivatives of formula (I) and the physiologically acceptable salts thereof, and the solvates thereof and the hydrates thereof; and a use of a substance selected from the group consisting of the pyrimidone derivatives of formula (I) and the physiologically acceptable salts thereof, and the solvates thereof and the hydrates thereof for the manufacture of
25 the aforementioned medicament.

As used herein, the C₁₋₆ alkyl group represents a straight or branched alkyl group having 1 to 6 carbon atoms, for example, methyl group, ethyl group, n-propyl group, isopropyl group, n-butyl group, isobutyl group, sec-butyl group, tert-butyl group, n-pentyl group, isopentyl group, neopentyl group, 1,1-dimethylpropyl group, n-hexyl group, isohexyl group, and the like;

The C₁₋₁₀ alkyl group represents a straight or branched alkyl group having 1 to 10 carbon atoms, for example in addition to the C₁₋₆ alkyl groups cited above,

- heptyl group, octyl group, nonyl group, decyl group, and the like;
- The C₂₋₆ alkylene group represents a divalent alkyl group;
- The C₂₋₆ alkenyl group represents an alkyl group having 2 to 6 carbon atoms and one or two double bond;
- 5 The C₃₋₆ alkynyl group represents an alkyl group having 3 to 6 carbon atoms and one or two triple bond;
- The C₁₋₆ alkoxy group represents an alkyloxy group having 1 to 6 carbon atoms for example, methoxy group, ethoxy group, propoxy group, isopropoxy group, butoxy group, isobutoxy group, sec-butoxy group, tert-butoxy group,
- 10 pentyloxy group, isopentyloxy group, neopentyloxy group, 1,1-dimethylpropyloxy group and the like;
- The C₁₋₆ acyloxy group represents an alkylcarbonyloxy group having 1 to 6 carbon atoms for example, methylcarbonyloxy group, ethylcarbonyloxy group, propylcarbonyloxy group, isopropylcarbonyloxy group, butylcarbonyloxy group,
- 15 isobutylcarbonyloxy group, sec-butylcarbonyloxy group, tert-butylcarbonyloxy group, pentylcarbonyloxy group, isopentylcarbonyloxy group, neopentylcarbonyloxy group, 1,1-dimethylpropylcarbonyloxy group and the like;
- The halogen atom represents a fluorine, chlorine, bromine or iodine atom;
- The C₁₋₂ perhalogenated alkyl group represents an alkyl group wherein all
- 20 the hydrogen have been substituted by a halogen atom, for example a CF₃ or C₂F₅;
- The C₁₋₃ halogenated alkyl group represents an alkyl group wherein at least one hydrogen has not been substituted by a halogen atom;
- 25 The C₆₋₁₀ aryl group represents a phenyl group, a naphth-1-yl group or a naphth-2-yl group;
- The C₆₋₁₀ ARYL group for R2 represents an indan-1-yl ring, an indan-2-yl ring tetrahydronaphthalen-1-yl ring, tetrahydronaphthalen-2-yl ring, a phenyl group, naphth-1-yl group or a naphth-2-yl group;
- 30 The C_{6,10} aryloxy group represents a phenoxy group, a 1-naphthyoxy group or a 2-naphthyoxy group;
- The C₁₋₆ monoalkylamino group represents an amino group substituted by one C₁₋₆ alky group, for example, methylamino group, ethylamino group, propylamino group, isopropylamino group, butylamino group, isobutylamino group, tert-butylamino group, pentylamino group and isopentylamino group;
- 35 The C₂₋₁₂ dialkylamino group represents an amino group substituted by two C₁₋₆ alkyl groups, for example, dimethylamino group, ethylmethylamino group, diethylamino group, methylpropylamino group and diisopropylamino group;
- The heterocyclic ring having 1-4 hetero atoms selected from oxygen atom,

sulfur atom, and nitrogen atom, and having total ring-constituting atoms of 5-10 represents, for example, a furan ring, dihydrofuran ring, tetrahydrofuran ring, pyran ring, dihydropyran ring, tetrahydropyran ring, benzofuran ring, furopyridine ring, isobenzofuran ring, chromene ring, chroman ring, isochroman ring, thiophene ring, 5 benzothiophene ring, thienopyridine ring, pyrrole ring, pyrrolidine ring, pyrrolidone ring, imidazole ring, imidazoline ring, imidazolidine ring, imidazopyridine ring, pyrazole ring, pyrazoline ring, pyrazolidine ring, triazole ring, tetrazole ring, pyridine ring, pyridine oxide ring, piperidine ring, pyrazine ring, piperazine ring, pyrimidine ring, pyridazine ring, indolizine ring, indole ring, indoline ring, isoindole 10 ring, isoindoline ring, indazole ring, benzimidazole ring, purine ring, quinolizine ring, quinoline ring, isoquinoline ring, phthalazine ring, naphtyridine ring, quinoxaline ring, quinazoline ring, cinnoline ring, pteridine ring, oxazole ring, oxazolidine ring, isoxazole ring, isoxazolidine ring, thiazole ring, benzothiazole ring, thiazylidine ring, isothiazole ring, isothiazolidine ring, dioxane ring, dithian 15 ring, morpholine ring, thiomorpholine ring, phthalimide ring, tetrahydropyridoindole ring, tetrahydroisoquinoline ring, tetrahydrothienopyridine ring, tetrahydrobenzofuropyridine ring, and the like.

When R2 represents a C₁₋₁₀ alkyl group, a C₂₋₆ alkenyl group, a C₃₋₆ alkynyl, a C₃₋₆ cycloalkyl group which may be substituted, these groups may have 20 1 or 3 substituents selected from the group consisting of a C₃₋₆ cycloalkyl, an adamantyl, a C₃₋₆ cycloalkyloxy group, a C₁₋₆ alkoxy group, a C₆₋₁₀ aryloxy group which may be substituted, a hydroxyl group, a C₁₋₆ alkylthio group, a C₆₋₁₀ arylthio group, an amino, a C₁₋₆ monoalkylamino group, a C₂₋₁₂ dialkylamino group, a C₆₋₁₀ 25 arylamino group, a C₁₋₆ acyloxy, a C₆₋₁₀ aryl group which may be substituted, a heterocyclic ring having 1-4 hetero atoms selected from oxygen atom, sulfur atom, and nitrogen atom, and having total ring-constituting atoms of 5-10 which may be substituted.

When a C₆₋₁₀ aryl group may be substituted, the C₆₋₁₀ aryl group may have 30 1 to 3 substituents selected from the group consisting of a C₁₋₆ alkyl group, halogen atom, a C₁₋₂ perhalogenated alkyl group, a C₁₋₃ halogenated alkyl group, a hydroxyl group, a C₁₋₆ alkoxy group, methylenedioxy group, a nitro, a cyano, an amino, a C₁₋₆ monoalkylamino group, a C₂₋₁₂ dialkylamino group, a (C₁₋₆- 35 alkyl)carbonylamino group, a (C₆₋₁₀-aryl)carbonylamino group, a (C₁₋₆-alkoxy)carbonylamino group, aminocarbonyl group, a (C₁₋₆-monoalkylamino)carbonyl group, a (C₂₋₁₂-dialkylamino)carbonyl group, a formyl, a C₁₋₆ alkylcarbonyl group, a (C₆₋₁₀-aryl)carbonyl group, a C₁₋₆ alkylsulfonyl group, a

$C_{6,10}$ arylsulfonyl group, aminosulfonyl group, a C_{1-6} monoalkylaminosulfonyl group, a C_{2-12} dialkylaminosulfonyl group, or a phenyl group;

Wherein the C_{1-6} alkyl groups and the C_{1-6} alkoxy groups are optionally substituted by a halogen atom, a hydroxyl group, a C_{1-6} alkoxy group, a C_{1-6} acyloxy group, an amino, a C_{1-6} monoalkylamino group, a C_{2-12} dialkylamino group, a (C_{1-6} -alkyl)carbonylamino group, an amino- C_{1-6} alkyl, a C_{1-6} monoalkylamino- C_{1-6} alkyl group, a C_{2-12} dialkylamino- C_{1-6} alkyl group, a (C_{1-6} -alkyl)carbonylamino group a ($C_{6,10}$ -aryl)carbonylamino group, a (C_{1-6} -alkoxy)carbonylamino group, a C_{1-6} alkylsulfonylamino group, a $C_{6,10}$ arylsulfonylamino group, a phenyl group or a heterocyclic ring having 1-4 hetero atoms selected from oxygen atom, sulfur atom, and nitrogen atom, and having total ring-constituting atoms of 5-10 which may be substituted.

When a $C_{6,10}$ ARYL group may be substituted, the $C_{6,10}$ ARYL group may have 1 to 3 substituents selected from the group consisting of a C_{1-6} alkyl group, halogen atom, a C_{1-2} perhalogenated alkyl group, a C_{1-3} halogenated alkyl group, a hydroxyl group, a C_{1-6} alkoxy group, methylenedioxy group, a nitro, a cyano, an amino, a C_{1-6} monoalkylamino group, a C_{2-12} dialkylamino group, a (C_{1-6} alkyl)carbonylamino group, a ($C_{6,10}$ aryl)carbonylamino group, a (C_{1-6} alkoxy)carbonylamino group.

When a $C_{6,10}$ aryloxy group may be substituted, the $C_{6,10}$ aryl group may have 1 to 3 substituents as defined above for the $C_{6,10}$ aryl group.

When the heterocyclic ring having 1-4 hetero atoms selected from oxygen atom, sulfur atom, and nitrogen atom, and having a total ring-constituting atoms of 5-10, may be substituted, the heterocyclic ring may have 1 to 3 substituents selected from the group consisting of a C_{1-6} alkyl group, halogen atom, a C_{1-2} perhalogenated alkyl group, a C_{1-3} halogenated alkyl group, a hydroxyl group, a C_{1-6} alkoxy group, a nitro, a cyano, an amino, a C_{1-6} monoalkylamino group, a C_{2-12} dialkylamino group, a (C_{1-6} -alkyl)carbonylamino group, a ($C_{6,10}$ aryl)carbonylamino group, a (C_{1-6} alkoxy)carbonylamino group, aminocarbonyl group, a (C_{1-6} monoalkylamino)carbonyl group, a (C_{2-12} dialkylamino)carbonyl group, a formyl, a C_{1-6} alkylcarbonyl group, a ($C_{6,10}$ aryl)carbonyl group, a C_{1-6} alkylsulfonyl group, a $C_{6,10}$ arylsulfonyl group, aminosulfonyl group, a C_{1-6} monoalkylaminosulfonyl group, a C_{2-12} dialkylaminosulfonyl group, or a phenyl group;

Wherein the C_{1-6} alkyl groups and the C_{1-6} alkoxy group being optionally

substituted by a halogen atom, a hydroxyl group, a C₁₋₆ alkoxy group, an amino, a C₁₋₆ monoalkylamino group, a C₂₋₁₂ dialkylamino group, a (C₁₋₆ alkyl)carbonylamino group, a (C_{6,10} aryl)carbonylamino group, a (C₁₋₆ alkoxy)carbonylamino group, a C₁₋₆ alkylsulfonylamino group, a C_{6,10} arylsulfonylamino group or a phenyl group.

When the C₂₋₆ alkylene group may be substituted, the C₂₋₆ alkylene group may have 1 to 3 substituents selected from a group consisting of a C₁₋₆ alkyl group which may be substituted by a C_{6,10} aryl group which may be substituted, a C₁₋₆ alkyl group which may be substituted by a heterocyclic ring which may be substituted, a C_{6,10} aryl group which may be substituted, a heterocyclic ring which may be substituted; the substituents being as defined here above.

When R1 and R2 form together a chain of formula -(CH₂)₂-X-(CH₂)₂- or -(CH₂)₂-X-(CH₂)₃- wherein X represents a nitrogen atom which may be substituted, the group NR1R2 represents a piperazine ring or homopiperazine which may be substituted in position 4 by a substituent selected from the group consisting of a C₁₋₆ alkyl group which may be substituted by a C_{6,10} aryl group which may be substituted or by a heterocyclic ring which may be substituted; a C_{6,10} aryl group which may be substituted or a heterocyclic ring which may be substituted, the substituents being as defined hereabove.

The compounds represented by the aforementioned formula (I) may form a salt. Examples of the salt include, when an acidic group exists, salts of alkali metals and alkaline earth metals such as lithium, sodium, potassium, magnesium, and calcium; salts of ammonia and amines such as methylamine, dimethylamine, trimethylamine, dicyclohexylamine, tris(hydroxymethyl)aminomethane, N,N-bis(hydroxyethyl)piperazine, 2-amino-2-methyl-1-propanol, ethanolamine, N-methylglucamine, and L-glucamine; or salts with basic amino acids such as lysine, δ-hydroxylysine, and arginine. The base-addition salts of acidic compounds are prepared by standard procedures well known in the art.

When a basic group exists, examples include salts with mineral acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid; salts with organic acids such as methanesulfonic acid, benzenesulfonic acid, p-toluenesulfonic acid, acetic acid, propionic acid, tartaric acid, fumaric acid, maleic acid, malic acid, oxalic acid, succinic acid, citric acid, benzoic acid, mandelic acid, cinnamic acid, lactic acid, glycolic acid, glucuronic acid, ascorbic acid, nicotinic

acid, and salicylic acid; or salts with acidic amino acids such as aspartic acid, and glutamic acid.

The acid-addition salts of the basic compounds are prepared by standard procedures well known in the art which include, but are not limited thereto, dissolving the free base in an aqueous alcohol solution containing the appropriate acid and isolating the salt by evaporating the solution, or by reacting the free base and an acid in an organic solvent, in which case the salt separates directly, or is precipitated with a second organic solvent, or can be obtained by concentration of the solution. The acids which can be used to prepare the acid-addition salts include preferably those which produce, when combined with the free base, pharmaceutically-acceptable salts, that is, salts whose anions are relatively innocuous to the animal organism in pharmaceutical doses of the salts, so that the beneficial properties inherent in the free base are not compromised by side effects ascribable to the anions. Although medicinally acceptable salts of the basic compounds are preferred, all acid-addition salts are within the scope of the present invention.

In addition to the pyrimidone derivatives represented by the aforementioned formula (I) and salts thereof, their solvates and hydrates also fall within the scope of the present invention. The pyrimidone derivatives represented by the aforementioned formula (I) may have one or more asymmetric carbon atoms. As for the stereochemistry of such asymmetric carbon atoms, they may independently be in either (R) and (S) configuration, and the pyrimidone derivative may exist as stereoisomers such as optical isomers, or diastereoisomers. Any stereoisomers in pure form, any mixtures of stereoisomers, racemates and the like fall within the scope of the present invention.

Examples of preferred compounds of the present invention are shown in table 1 hereinafter. However, the scope of the present invention is not limited by these compounds.

Preferred compounds of the present invention represented by formula (I) include also:

35

- (1) Compounds wherein R3 represents a 3- or 4-pyridyl group and more preferably 4-pyridyl group which may be substituted by a C₁₋₂ alkyl group, C₁₋₂ alkoxy group or halogen atom;

- (2) Compounds wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group which may be substituted by a phenyl group;
- (3) Compounds wherein R2 represents an unsubstituted C₁₋₁₀ alkyl group;
- (4) Compounds wherein R2 represents a substituted C₁₋₆ alkyl group, a C₃₋₆ cycloalkyl group which may be substituted, an indanyl group which may be substituted or a C₂₋₄ alkenyl group which may be substituted;
- 5 (5) Compounds wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group and R2 represents a C₁₋₆ alkyl group which may be substituted, a C₃₋₆ cycloalkyl group which may be substituted, an indanyl group which may be substituted;
- 10 (6) Compounds wherein R1 and R2 form together a C₂₋₆ alkylene group.

More preferred compounds of the present invention represented by formula (I) include also:

- (1) Compounds wherein R3 represents an unsubstituted 4-pyridyl group;
- 15 (2) Compounds wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group which may be substituted by a phenyl group and R2 represents a C₁₋₆ alkyl group which may be substituted by a C₃₋₆ cycloalkyl, an adamantyl, a C₁₋₆ alkoxy group, a hydroxyl group, a phenylthio group, an amino, a C₁₋₆ monoalkylamino group, a C₂₋₁₂ dialkylamino group, a phenylamino group, a C₁₋₆ acyloxy, a phenyl group which may be substituted, a heterocyclic ring having 1-4 hetero atoms selected from oxygen atom, sulfur atom, and nitrogen atom, and having total ring-constituting atoms of 5-10 which may be substituted, preferably the heterocyclic ring being selected from an indole ring or a substituted indole ring, a thiophene or substituted thiophene ring, a pyridine ring and a piperidine ring;
- 20 (3) Compounds wherein R3 is defined as specified under (1), and R1 and R2 are specified under (2) for the more preferred compounds;
- (4) Compounds wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group and R2 represents an indanyl group or an substituted indanyl group.
- 25

30 Particularly preferred compounds of the present invention represented by formula (I) include:

- 2-[[2-(phenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
35 2-[[2-(3-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(3-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

- 2-[[2-(4-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-bromophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2,4-dichlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
5 2-[[2-(4-aminophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(3,4-dimethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2,5-dimethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[2-(4-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
10 2-[[2-(4-hydroxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-methylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one
2-[[2-(4-aminosulfonylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[2-(3-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
15 2-[[2-(thiophen-2-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[4-(phenyl)butyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-phenylmethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[2-(4-phenylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
20 2-[(phenylmethyl)amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-methoxyphenyl)methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2,5-dimethoxyphenyl)ethyl]methylamino]-3-methyl-6-pyridin-4-ylpyrimidin-
4(3H)-one,
2-[[3-(3-aminopropoxy)phenyl]methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-
25 4(3H)-one,
2-[[3-(aminomethyl)phenyl]methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[3-(phenyl)propyl]amino]-3-phenylmethyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
30 2-[[2-(5-methoxy-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[2-(5-phenylmethoxy-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-
ylpyrimidin-4(3H)-one,
2-[[2-(7-methyl-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
35 one,
2-[[2-(1-methyl-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
one,
2-[[2-(1-methyl-1H-indol-3-yl)ethyl]methylamino]-3-methyl-6-pyridin-4-ylpyrimidin-

- 4(3H)-one,
2-(cyclopentylamino)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(ethylamino)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[(indan-2-yl)amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
5 2-(piperidin-1-yl)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(pyrrolidin-1-yl)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(5-Amino-pentylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(4-Amino-butylamino)-3-(3-phenyl-propyl)-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(6-Amino-hexylamino)-3-(3-phenyl-propyl)-6-pyridin-4-yl-3H-pyrimidin-4-one,
10 2-(5-Amino-pentylamino)-3-phenethyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(6-Amino-hexylamino)-3-phenethyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(4-Amino-butylamino)-3-phenethyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-Cyclohexylamino-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-Butylamino-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
15 3-Methyl-2-pentylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-Hexylamino-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-Heptylamino-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-octylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-nonylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
20 2-Decylamino-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(2-Cyclohexyl-ethylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(3-methyl-butylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(Cyclohexylmethyl-amino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(2-propoxy-ethylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
25 2-(3-Cyclohexyl-propylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(3-Ethoxy-propylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-[(5-Amino-pentyl)-phenetyl-amino]-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(5-Hydroxy-pentylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(4-Hydroxy-butylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
30 2-(3-Isopropoxy-propylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(3-propoxy-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(2-Hydroxy-2-phenyl-ethylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(2-Cyclopentyl-ethylamino)-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(3-piperidin-1-yl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
35 2-[(3-Cyclohexyl-propyl)-methyl-amino]-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
Acetic acid 2-(1-methyl-6-oxo-4-pyridin-4-yl-1,6-dihydro-pyrimidin-2-ylamino)-1-phenyl-ethyl ester

- 2-(2-Adamantan-1-yl-ethylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-[3-(3-Hydroxy-propoxy)-benzylamino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-methyl-2-[3-(2-piperidin-4-yl-ethoxy)-benzylamino]-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
5 Acetic acid 3-{3-[(1-methyl-6-oxo-4-pyridin-4-yl-1,6-dihydro-pyrimidin-2-ylamino)-methyl]-phenoxy}-propyl ester
2-[3-(3-Dimethylamino-propoxy)-benzylamino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
10 3-Methyl-2-{methyl-[2-(4-methylaminomethyl-phenyl)-ethyl]-amino}-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-[(4-Amino-butyl)-phenethyl-amino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(3-Amino-propyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
15 3-(5-Amino-pentyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(4-Amino-butyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,

3-Methyl-6-pyridin-4-yl-2-[3-(3-pyridin-4-yl-propoxy)-benzylamino]-3*H*-pyrimidin-4-one,
20 3-(6-Amino-hexyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(6-Amino-hexyl)-2-phenethylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(2-Amino-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(3-Amino-propyl)-2-[2-(2-methoxy-phenyl)-ethylamino]-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
25 3-(2-Hydroxy-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(2-Dimethylamino-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(5-Amino-pentyl)-2-phenethylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-(4-Amino-butyl)-2-phenethylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
30 2-(4-Amino-butylamino)-3-benzyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(6-Amino-hexylamino)-3-benzyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(2-phenylthio-ethylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(2-phenylamino-ethylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(2-phenoxy-ethylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
35 3-Methyl-2-(3-phenyl-allylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one, and
3-Methyl-6-pyridin-4-yl-2-(2-pyridin-2-yl-ethylamino)-3*H*-pyrimidin-4-one.

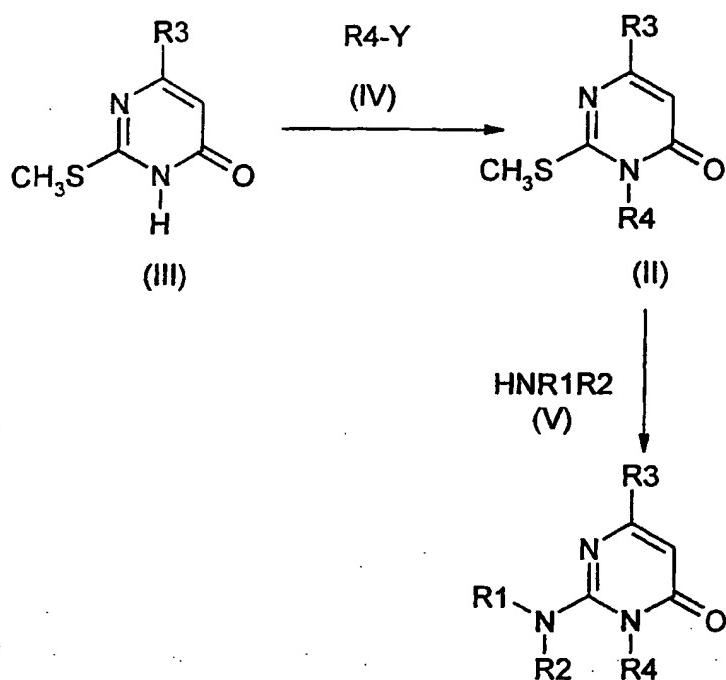
As a further object, the present invention concerns also methods for preparing the pyrimidone compounds represented by the aforementioned formula (I).

These compounds can be prepared, for example, according to the methods
5 explained below.

1. Preparation Method 1

Pyrimidone compounds represented by the aforementioned formula (I)
10 may be prepared according to scheme 1.

Scheme 1



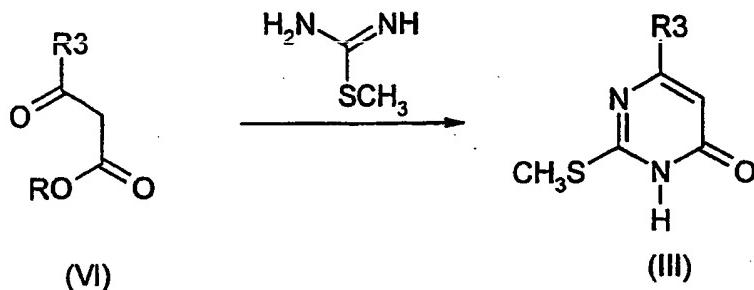
15 (In the above scheme the definition of R1, R2, R3 and R4 are the same as those already described above for compounds of formula (I)).

The 2-methylthio derivative represented by the above formula (III), wherein R3 is as defined for compound of formula (I), is allowed to react with a
20 compound of formula (IV), wherein Y represents a halogen atom such as for example a bromine or iodine in the presence of a base such as for example potassium carbonate, to obtain a compound of formula (II). The reaction may be

carried out in aprotic polar solvents such as formamide, N,N-dimethylformamide, N,N-dimethylacetamide, N-methylpyrrolidone and the like, at a suitable temperature ranging from -10 to + 20 °C under ordinary air.

- 5 Compound of formula (II) may then react with an amine of formula (V) to obtain the compound of the aforementioned formula (I). The reaction may be carried out in pyridine in presence of 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU), at a suitable temperature ranging from 25°C to reflux temperature.
- 10 Compound of formula (III) may be prepared according to the method defined in scheme 2.

Scheme 2



- 15 (In the above scheme R represents an alkyl group and the definition of R2 and R3 are the same as those already described for compound of formula (I).)

- 20 According to this method, the 3-ketoester of formula (VI) is allowed to react with a 2-methyl-2-thiopseudourea sulfate in the presence of a base such as potassium hydroxide. The reaction may be carried out in solvent such as water or an alcohol, such as ethanol, propanol and butanol, at a suitable temperature ranging from 25-100°C under ordinary air.

- 25 Compounds of formula (IV), (V) and formula (VI) are commercially available or may be synthesized according to known methods of one skilled in the art.

- 30 For example compounds of formula (VI), wherein R, R2 and R3 are as defined above, can be prepared by reacting a nicotinic acid optionally substituted

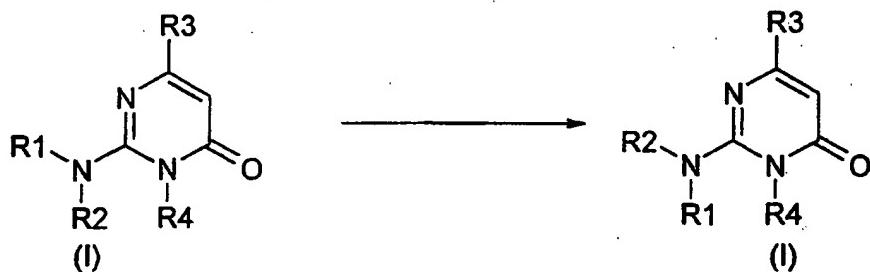
by a C₁₋₄ alkyl group, C₁₋₄ alkoxy group or an halogen, with a malonic acid monoester. The reaction can be carried out using methods well known to one skilled in the art, such as for example in presence of a coupling agent such as 1,1'-carbonylbis-1H-imidazole in a solvent such as a tetrahydrofuran at a temperature ranging from 20 to 70°C.

2. Preparation method 2

Alternatively pyrimidone compounds represented by the aforementioned formula (I) may be prepared according to scheme 2.

Scheme 2

15



Wherein R1 = H

Wherein R1 =
optionally substituted alkyl group

20

Compounds of formula (I) wherein R1 represents a hydrogen atom, can be alkylated by methods well known to one skilled in the art such as, for example, by reacting (I), wherein R1 represents a hydrogen atom, with sodium hydride, in an aprotic polar such as dimethylacetamide or dimethylformamide at a temperature ranging from 0° to 10°. An alkylating agent such as an optionally substituted C₁₋₆ alkyl halide is then added to obtain the compound of the above mentioned formula (I) wherein R1 represents an optionally substituted C₁₋₆ alkyl group.

25

In addition when applicable, compound of formula (I) can be derivatised affording other compounds of formula (I), using well known methods in the art, for example when the C₆₋₁₀ aryl groups or the heterocyclic ring is substituted by a hydroxyl group, the hydroxyl group can be alkylated to give a C₁₋₈ alkoxy group, or when the C₆₋₁₀ ARYL group, the C₆₋₁₀ aryl group or the heterocyclic ring is substituted by an amino group or an aminoalkyl group, the amino function can be

alkylated, acylated, etc... to give the corresponding derivatives.

In the above reactions, protection or deprotection of a functional group may sometimes be necessary. A suitable protecting group can be chosen 5 depending on the type of a functional group, and a method described in the literature may be applied. Examples of protecting groups, of protection and deprotection methods are given for example in *Protective groups in Organic Synthesis* Greene et al., 2nd Ed. (John Wiley & Sons, Inc., New York).

10 The compounds of the present invention have inhibitory activity against GSK3 β . Accordingly, the compounds of the present invention are useful as an active ingredient for the preparation of a medicament, which enables preventive and/or therapeutic treatment of neurodegenerative diseases such as Alzheimer's disease. In addition, the compounds of the present invention are also useful as an 15 active ingredient for the preparation of a medicament for preventive and/or therapeutic treatment of neurodegenerative diseases such as Parkinson's disease, tauopathies (e.g. frontotemporoparietal dementia, corticobasal degeneration, Pick's disease, progressive supranuclear palsy) and other dementia including vascular dementia; acute stroke and others traumatic injuries; cerebrovascular 20 accidents (e.g. age related macular degeneration); brain and spinal cord trauma; peripheral neuropathies; retinopathies and glaucoma; and other diseases such as non-insulin dependent diabetes (such as diabetes type II) and obesity; manic depressive illness; schizophrenia; alopecia; cancers such as breast cancer, non-small cell lung carcinoma, thyroid cancer, T or B-cell leukemia and several virus- 25 induced tumors.

The present invention further relates to a method for treating neurodegenerative diseases caused by abnormal activity of GSK3 β and of the aforementioned diseases which comprises administering to a mammalian 30 organism in need thereof an effective amount of a compound of the formula (I).

As the active ingredient of the medicament of the present invention, a substance may be used which is selected from the group consisting of the compound represented by the aforementioned formula (I) and pharmacologically 35 acceptable salts thereof, and solvates thereof and hydrates thereof. The substance, per se, may be administered as the medicament of the present invention, however, it is desirable to administer the medicament in a form of a pharmaceutical composition which comprises the aforementioned substance as an

active ingredient and one or more of pharmaceutical additives. As the active ingredient of the medicament of the present invention, two or more of the aforementioned substances may be used in combination. The above pharmaceutical composition may be supplemented with an active ingredient of 5 another medicament for the treatment of the above mentioned diseases. A type of the pharmaceutical composition is not particularly limited, and the composition may be provided as any formulation for oral or parenteral administration. For example, the pharmaceutical composition may be formulated, for example, in the form of pharmaceutical compositions for oral administration such as granules, fine 10 granules, powders, hard capsules, soft capsules, syrups, emulsions, suspensions, solutions and the like, or in the form of pharmaceutical compositions for parenteral administrations such as injections for intravenous, intramuscular, or subcutaneous administration, drip infusions, transdermal preparations, transmucosal preparations, nasal drops, inhalants, suppositories and the like. Injections or drip 15 infusions may be prepared as powdery preparations such as in the form of lyophilized preparations, and may be used by dissolving just before use in an appropriate aqueous medium such as physiological saline. Sustained-release preparations such as those coated with a polymer may be directly administered intracerebrally.

20

Types of pharmaceutical additives used for the manufacture of the pharmaceutical composition, content ratios of the pharmaceutical additives relative to the active ingredient, and methods for preparing the pharmaceutical composition may be appropriately chosen by those skilled in the art. Inorganic or 25 organic substances, or solid or liquid substances may be used as pharmaceutical additives. Generally, the pharmaceutical additives may be incorporated in a ratio ranging from 1% by weight to 90% by weight based on the weight of an active ingredient.

Examples of excipients used for the preparation of solid pharmaceutical 30 compositions include, for example, lactose, sucrose, starch, talc, cellulose, dextrin, kaolin, calcium carbonate and the like. For the preparation of liquid compositions for oral administration, a conventional inert diluent such as water or a vegetable oil may be used. The liquid composition may contain, in addition to the inert diluent, auxiliaries such as moistening agents, suspension aids, sweeteners, aromatics, 35 colorants, and preservatives. The liquid composition may be filled in capsules made of an absorbable material such as gelatin. Examples of solvents or suspension mediums used for the preparation of compositions for parenteral administration, e.g. injections, suppositories, include water, propylene glycol,

polyethylene glycol, benzyl alcohol, ethyl oleate, lecithin and the like. Examples of base materials used for suppositories include, for example, cacao butter, emulsified cacao butter, lauric lipid, witepsol.

5 Dose and frequency of administration of the medicament of the present invention are not particularly limited, and they may be appropriately chosen depending on conditions such as a purpose of preventive and/or therapeutic treatment, a type of a disease, the body weight or age of a patient, severity of a disease and the like. Generally, a daily dose for oral administration to an adult may 10 be 0.01 to 1,000 mg (the weight of an active ingredient), and the dose may be administered once a day or several times a day as divided portions, or once in several days. When the medicament is used as an injection, administrations may preferably be performed continuously or intermittently in a daily dose of 0.001 to 100 mg (the weight of an active ingredient) to an adult.

15

Chemical Examples

The present invention will be explained more specifically with reference to the following general examples, however, the scope of the present invention is not 20 limited to these examples.

Example 1 : Preparation of substituted 2-amino-3-methylpyrimidinones (method 1)

1.1. Preparation of Ethyl 3-(4-pyridyl)-3-oxopropionate

25

Isonicotinic acid (35.56 g, 289 mmol) was added to a solution of 1,1'-carbonylbis-1H-imidazole (46.98 g, 290 mmol) in tetrahydrofuran (700ml), and the resulting solution was stirred for 1.5 hr at 50°C. After cooling to room temperature, malonic acid monoester potassium salt (51.7 g, 304 mmol) and magnesium chloride (34.33 g, 361 mmol) were added, and the mixture was refluxed for 1 hr and then heated 30 at 50°C for 6 hr. The solvent was removed under reduced pressure and the residue was quenched by the addition of dilute acetic acid. The organic layer was extracted with ethyl acetate (3 times) and the combined extracts were washed with dilute aqueous sodium bicarbonate and brine, and were concentrated under 35 reduced pressure. The residue was purified by silica gel column chromatography (eluent ; hexane/ethyl acetate = 2/1 to 1/1) and recrystallization from hexane - ethyl acetate gave 41.52 g (74%) of the title compound.

1.2. Preparation of 2-(Methylthio)-6-pyridinyl-4-ylpyrimidin-4(1H)-one

To a solution of 5.76 g (20.7 mmol) of 2-methyl-2-thiopseudoiourea sulfate in 48 ml of water was added 4.85 g (86.52 mmol) of potassium hydroxide. The mixture was agitated and 8.0 g (41.4 mmol) of ethyl 3-(4-pyridyl)-3-oxopropionate was added and stirring was maintained for 48 hours.

The precipitate was recovered by filtration and was washed with water and then ether. The product was dried at 90°C in vacuo to give 6.26 g, 69% of white solid.
Mp : 328-330°C.

10

1.3. Preparation of 3-methyl-2-(methylthio)-6-pyridin-4-yl-pyrimidin-4(3H)-one

To 3.0 g (13.7 mmol) of 2-methylthio-6-(4-pyridyl)pyrimidin-4-one in 50 ml of dimethylformamide was added 2.08 g (15.05 mmol) of potassium carbonate, followed by 0.85 ml (13.68 mmol) of methyl iodide at 0°C and stirring was maintained for 1.5 hours.

The reaction mixture was added to cold water and extracted with dichloromethane. The solvent was evaporated and the resulting solid was purified by chromatography on silica gel, eluting with a mixture of dichloromethane/methanol (99:1 to 90:10) to give 2.36 g, 78% of a white solid.
Mp. 176-178°C.

1.4. Preparation of substituted 2-amino-3-methylpyrimidinones

25 A solution of 1 equivalent of 3-methyl-2-(methylthio)-6-pyridin-4-yl-pyrimidin-(3H)-one and 1-5 equivalents of an amine of formula HNR4R5 were suspended in pyridine (0.1-1M) containing 3 equivalents of the DBU (1,8-diazabicyclo[5.4.0]undec-7-ene) and was refluxed 24 hours.

The cooled solution was treated with a saturated aqueous solution of ammonium chloride and extracted with dichloromethane. The organic layer was dried and evaporated to give crude product which was purified by chromatography on silica gel.

2. Preparation of substituted 2-alkylamino-3-alkylpyrimidinones (method 2)

35

To a cooled (0°C) solution of substituted 2-amino-3-methylpyrimidinone (1 equivalent, 0.1 mole) in N,N-dimethylacetamide (0.35 ml) was added sodium hydride (0.11 mmole). The mixture was stirred for 5 min and alkyl iodide (0.1

mmole) was added, stirred for further 20 min at 0°C and then for 40 min at room temperature. Water (10 ml) was added, and the reaction mixture was extracted with ethyl acetate (3x 3 ml). The organic phases were separated, dried over sodium sulfate and evaporated to afford a residue which was purified by chromatography 5 on silica gel.

A list of chemical structures and physical data for compounds of the aforementioned formula (I) illustrating the present invention is given in table 1. The compounds have been prepared according to the examples.

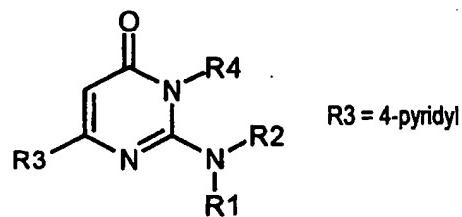
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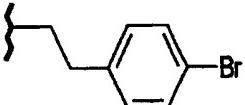
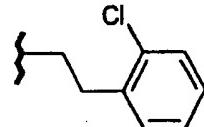
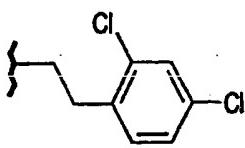
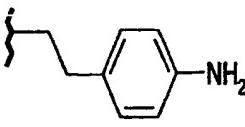
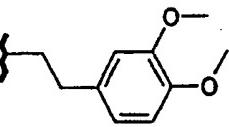
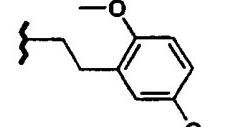
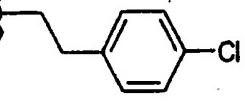
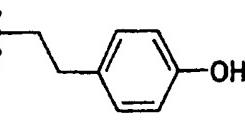
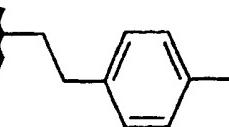
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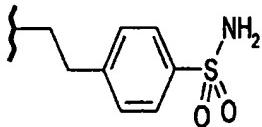
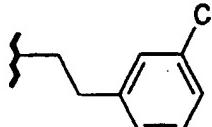
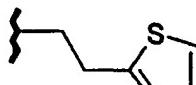
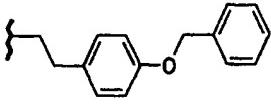
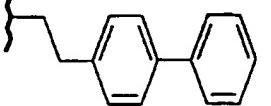
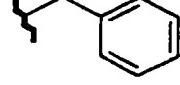
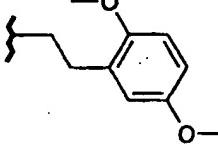
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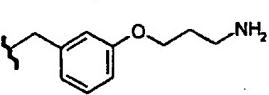
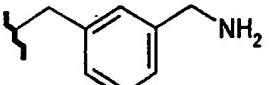
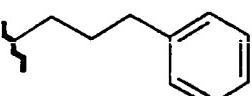
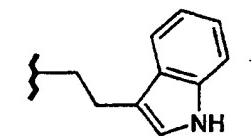
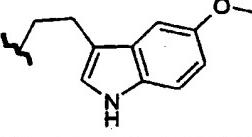
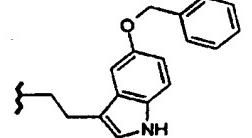
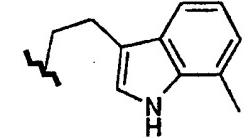
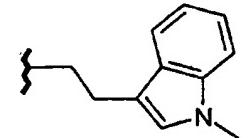
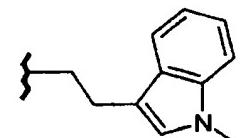
15



No.	R1	R2	R4	m.p. (°C)	[M+H] ⁺
1	H		Me	186-187.5	307
2	H		Me	142.4-142.6	337
3	H		Me		337
4	H		Me	149.2-149.5	337
5	H		Me	184.0-187.2	325
6	H		Me	158.9-159.2	325
7	H		Me	178.8-178.9	325

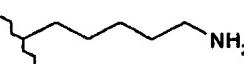
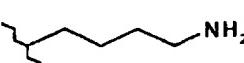
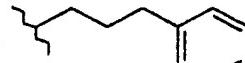
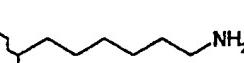
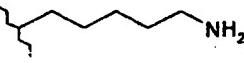
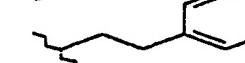
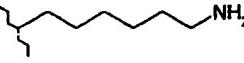
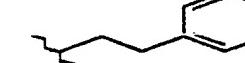
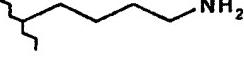
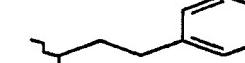
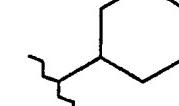
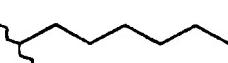
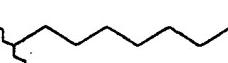
No.	R1	R2	R4	m.p. (°C)	[M+H] ⁺
8	H		Me	192.2-192.3	386
9	H		Me	175.2-175.4	341
10	H		Me	189.5-189.7	376
11	H		Me	197.3-197.5	322
12	H		Me	187.0-187.1	367
13	H		Me	180.9-181.3	367
14	H		Me	165.3-165.5	341
15	H		Me	257.4-257.9	323
16	H		Me	184.7-185	321

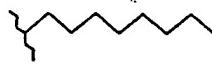
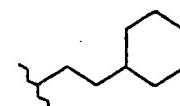
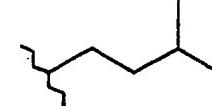
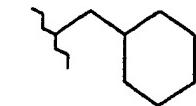
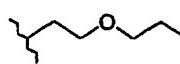
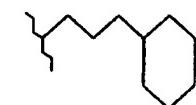
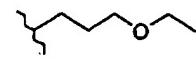
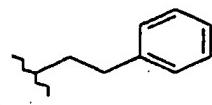
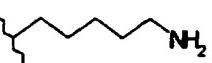
No.	R1	R2	R4	m.p. (°C)	[M+H] ⁺
17	H		Me	-	386
18	H		Me	162.8-163	341
19	H		Me	171.7-171.9	313
20	H		Me	-	335
21	H		Me	169.7-169.8	413
22	H		Me	175.4-175.6	383
23	H		Me	-	293
24	H		Me	-	323
25	Me		Me	-	381

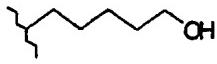
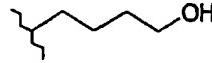
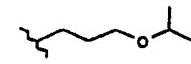
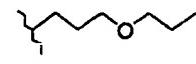
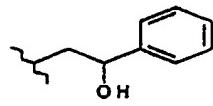
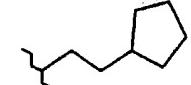
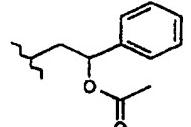
No.	R1	R2	R4	m.p. (°C)	[M+H] ⁺
26	H		Me	193-196 (*)	-
27	H		Me	193-197 (*)	-
28	H		CH ₂ -Ph	218-221	-
29	H		Me	217-218 (□)	346
30	H		Me	242.7-243.0	376
31	H		Me	168.4-168.6	452
32	H		Me	217.2-217.3	360
33	H		Me	-	360
34	Me		Me	-	374

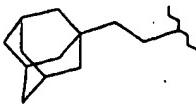
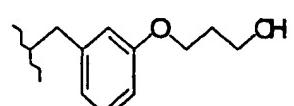
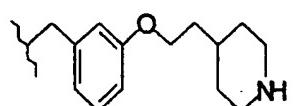
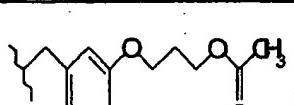
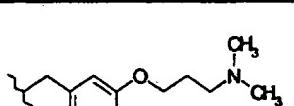
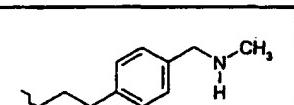
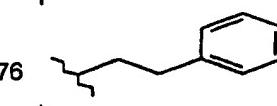
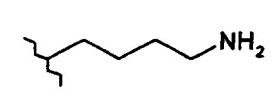
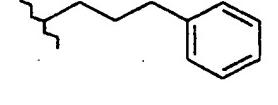
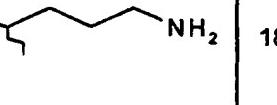
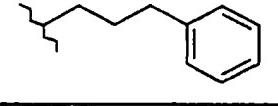
No.	R1	R2	R4	m.p. (°C)	[M+H] ⁺
35	H		Me	-	271
36	H	Et	Me	239-241 (□)	231
37	H		Me	211.5-211.8	319
38		- (CH ₂) ₅ -	Me	228-230 (***)	271
39		- (CH ₂) ₄ -	Me	-	257

All compounds are bases, except (*) : dihydrochloride, (**) : monohydrochloride,
 (□) : oxalate and (***): tartrate

40	H		Me	259-262(*)
41	H			134-136(*)
42	H			142-143(*)
43	H			235-238(*)
44	H			225-228(*)
45	H			246-248(*)
46	H		Me	268-270
47	H		Me	180-182
48	H		Me	157-159
49	H		Me	133-135
50	H		Me	121-123

51	H		Me	114-116
52	H		Me	113-115
53	H		Me	162-164(**)
54	H		Me	146-150
55	H		Me	213-217
56	H		Me	218-220
57	H		Me	147-150
58	H		Me	177-180
59	H		Me	118-120
60			Me	158-162(*)

61	H		Me	164-165
62	H		Me	159-161
63	H		Me	95-98
64	H		Me	107-109
65	H		Me	202-206
66	H		Me	190-194
67	H		Me	162-163
68	Me		Me	188-191(**)
69	H		Me	81-84

70	H		Me	107-111
71	H		Me	98-101
72	H		Me	225-232(*)
73	H		Me	198-201
74	H		Me	80-82
75	Me		Me	157-160(*)
76		 	Me	210-213(**)
77	H	 		185-189(*)
78	H		Me	122-124

79	H			123-126(*)
80	H			124-127(*)
81	H		Me	210-213(*)
82	H			192-195(*)
83	H			125-127(*)
84	H			145-148(*)
85	H			198-201(*)
86	H			143-145
87	H			212-215(*)

88	H			178-180(*)
89	H			170-175(*)
90	H			163-168dec(*)
91	H			292-298(*)
92	H		Me	162-168
93	H		Me	102-105
94	H		Me	193-194

95	H		Me	198-200(**)
96	H		Me	187-189(**)
97	H		Me	154-156dec

All compounds are bases, except (*) : dihydrochloride, (**) : monohydrochloride,
 (□) : oxalate and (***): tartrate

Test Example: Inhibitory activity of the medicament of the present invention against GSK3 β :

Two different protocols can be used.

5

In a first protocol : 7.5 μ M of prephosphorylated GS1 peptide and 10 μ M ATP (containing 300,000 cpm of 33P-ATP) were incubated in 25 mM Tris-HCl, pH 7.5, 0.6 mM DTT, 6 mM MgCl₂, 0.6 mM EGTA, 0.05 mg/ml BSA buffer for 1 hour at room temperature in the presence of GSK3 β (total reaction volume : 100 microliters).

10
In a second protocol : 4.1 μ M of prephosphorylated GS1 peptide and 42 μ M ATP (containing 260,000 cpm 33P-ATP) were incubated in 80 mM Mes-NaOH, pH 6.5, 1 mM Mg acetate, 0.5 mM EGTA, 5 mM 2-mercaptoethanol, 0.02% Tween 20, 15 10% glycerol buffer for 2 hours at room temperature in the presence of GSK3 β . Inhibitors were solubilised in DMSO (final solvent concentration in the reaction medium, 1%).

20
The reaction was stopped with 100 microliters of a solution made of 25 g polyphosphoric acid (85% P₂O₅), 126 ml 85% H₃PO₄, H₂O to 500 ml and then diluted to 1:100 before use. An aliquot of the reaction mixture was then transferred to Whatman P81 cation exchange filters and rinsed with the solution described above. Incorporated 33P radioactivity was determined by liquid scintillation spectrometry.
25
The phosphorylated GS-1 peptide had the following sequence : NH₂-YRRAAVPPSPSLSRHSSPHQS(P)EDEE-COOH.

30
The GSK3 β inhibitory activity of the compounds of the present invention are expressed in IC₅₀, and as an illustration the range of IC₅₀'s of the compounds in table 1 is between 0.1 to 10 micromolar concentrations.

Formulation Example

(1) Tablets

5 The ingredients below were mixed by an ordinary method and compressed by using a conventional apparatus.

	Compound of Example 1	30 mg
	Crystalline cellulose	60 mg
	Corn starch	100 mg
10	Lactose	200 mg
	Magnesium stearate	4 mg

(2) Soft capsules

The ingredients below were mixed by an ordinary method and filled in soft capsules.

	Compound of Example 1	30 mg
	Olive oil	300 mg
	Lecithin	20 mg

20 (1) Parenteral preparations

The ingredients below were mixed by an ordinary method to prepare injections contained in a 1 ml ampoule.

	Compound of Example 1	3 mg
25	Sodium chloride	4 mg
	Distilled water for injection	1 ml

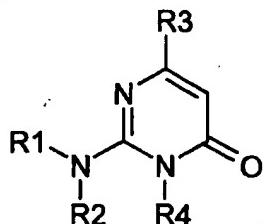
Industrial Applicability

30

The compounds of the present invention have GSK3 β inhibitory activity and are useful as an active ingredient of a medicament for preventive and/or therapeutic treatment of neurodegenerative diseases caused by abnormal activity of GSK3 β .

What is claimed is:

1. A pyrimidone derivative represented by formula (I) or a salt thereof, or a
 5 solvate thereof or a hydrate thereof:



(I)

Wherein :

- R1 represents a hydrogen atom or a C₁₋₆ alkyl group which may be substituted by a C_{6,10} aryl group;
- 10 R2 represents a C₁₋₁₀ alkyl group which may be substituted, a C₂₋₆ alkenyl group which may be substituted, a C₃₋₆ alkynyl group which may be substituted, a C₃₋₆ cycloalkyl group which may be substituted, or a C₆₋₁₀ ARYL group which may be substituted;
 or R1 and R2 form together a C₂₋₆ alkylene group which may be substituted;
- 15 or R1 and R2 form together a chain of formula -(CH₂)₂-X-(CH₂)₂- or -(CH₂)₂-X-(CH₂)₃- where X represents a oxygen atom, a sulfur atom, or a nitrogen atom which may be substituted;
- R3 represents a 2, 3 or 4-pyridyl group optionally substituted by a C₁₋₄ alkyl group, C₁₋₄ alkoxy group or halogen atom; and
- 20 R4 represents a C₁₋₁₀ alkyl group optionally substituted by a hydroxyl group, amino, C₁₋₆ monoalkylamino group, C₂₋₁₂ dialkylamino group or C_{6,10} aryl group which may be substituted.

2. A pyrimidone derivative or a salt thereof, or a solvate thereof or a
 25 hydrate thereof according to claim 1, wherein R3 represents an unsubstituted 4-pyridyl group.

3. A pyrimidone derivative or a salt thereof, or a solvate thereof or a
 30 hydrate thereof according to claim 1, wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group which may be substituted by a phenyl group.

4. A pyrimidone derivative or a salt thereof, or a solvate thereof or a hydrate thereof according to claim 1, wherein R2 represents an unsubstituted C₁₋₁₀ alkyl group.

5

5. A pyrimidone derivative or a salt thereof, or a solvate thereof or a hydrate thereof according to claim 1, wherein R2 represents a substituted C₁₋₆ alkyl group, a C₃₋₆ cycloalkyl group which may be substituted, an indanyl group which may be substituted or a C₂₋₄ alkenyl group which may be substituted.

10

6. A pyrimidone derivative or a salt thereof, or a solvate thereof or a hydrate thereof according to claim 1, wherein R1 and R2 form together a C₂₋₆ alkylene group.

15

7. A pyrimidone derivative or a salt thereof, or a solvate thereof or a hydrate thereof according to claim 1 or 2, wherein R1 represents a hydrogen atom or a C₁₋₃ alkyl group which may be substituted by a phenyl group and R2 represents a C₁₋₆ alkyl group which may be substituted by a C₃₋₆ cycloalkyl, an adamantyl, a C₁₋₆ alkoxy group, a hydroxyl group, a phenylthio group, an amino, a C₁₋₆ monoalkylamino group, a C₂₋₁₂ dialkylamino group, a phenylamino group, a C₁₋₆ acyloxy, a phenyl group which may be substituted, a heterocyclic ring being selected from an indole ring or substituted indole ring, a thiophene or substituted thiophene ring, a pyridine ring and a piperidine ring.

25

8. A pyrimidone derivative which is selected from the group consisting of:

2-[[2-(phenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(4-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(3-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(2-methoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

30

2-[[2-(2-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(3-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(4-fluorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(4-bromophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(2-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

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2-[[2-(2,4-dichlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(4-aminophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

2-[[2-(3,4-dimethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

- 2-[[2-(2,5-dimethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-hydroxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
5 2-[[2-(4-methylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one
2-[[2-(4-aminosulfonylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(3-chlorophenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(thiophen-2-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
10 2-[[4-(phenyl)butyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-phenylmethoxyphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(4-phenylphenyl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[(phenylmethyl)amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
15 2-[[2-(2,5-dimethoxyphenyl)methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(2,5-dimethoxyphenyl)ethyl]methylamino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[[3-(3-aminopropoxy)phenyl]methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
20 2-[[[3-(aminomethyl)phenyl]methyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[3-(phenyl)propyl]amino]-3-phenylmethyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(5-methoxy-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-
25 one,
2-[[2-(5-phenylmethoxy-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(7-methyl-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
30 2-[[2-(1-methyl-1H-indol-3-yl)ethyl]amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[[2-(1-methyl-1H-indol-3-yl)ethyl]methylamino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(cyclopentylamino)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
35 2-(ethylamino)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-[(indan-2-yl)amino]-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(piperidin-1-yl)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,
2-(pyrrolidin-1-yl)-3-methyl-6-pyridin-4-ylpyrimidin-4(3H)-one,

- 2-(5-Amino-pentylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(4-Amino-butylamino)-3-(3-phenyl-propyl)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(6-Amino-hexylamino)-3-(3-phenyl-propyl)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(5-Amino-pentylamino)-3-phenethyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
5 2-(6-Amino-hexylamino)-3-phenethyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(4-Amino-butylamino)-3-phenethyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-Cyclohexylamino-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-Butylamino-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-pentylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
10 2-Hexylamino-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-Heptylamino-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-octylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-nonylamino-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-Decylamino-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
15 2-(2-Cyclohexyl-ethylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(3-methyl-butylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(Cyclohexylmethyl-amino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(2-propoxy-ethylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(3-Cyclohexyl-propylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
20 2-(3-Ethoxy-propylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-[(5-Amino-pentyl)-phenetyl-amino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(5-Hydroxy-pentylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(4-Hydroxy-butylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(3-Isopropoxy-propylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
25 3-Methyl-2-(3-propoxy-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(2-Hydroxy-2-phenyl-ethylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-(2-Cyclopentyl-ethylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
3-Methyl-2-(3-piperidin-1-yl-propylamino)-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-[(3-Cyclohexyl-propyl)-methyl-amino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-
30 one,
Acetic acid 2-(1-methyl-6-oxo-4-pyridin-4-yl-1,6-dihydro-pyrimidin-2-ylamino)-1-phenyl-ethyl ester
2-(2-Adamantan-1-yl-ethylamino)-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
2-[3-(3-Hydroxy-propoxy)-benzylamino]-3-methyl-6-pyridin-4-yl-3*H*-pyrimidin-4-
35 one,
3-methyl-2-[3-(2-piperidin-4-yl-ethoxy)-benzylamino]-6-pyridin-4-yl-3*H*-pyrimidin-4-one,
Acetic acid 3-{3-[(1-methyl-6-oxo-4-pyridin-4-yl-1,6-dihydro-pyrimidin-2-ylamino)-

- methyl]-phenoxy}-propyl ester
2-[3-(3-Dimethylamino-propoxy)-benzylamino]-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-{methyl-[2-(4-methylaminomethyl-phenyl)-ethyl]-amino}-6-pyridin-4-yl-
5 3H-pyrimidin-4-one,
2-[(4-Amino-butyl)-phenethyl-amino]-3-methyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-(3-Amino-propyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-(5-Amino-pentyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
10 3-(4-Amino-butyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,

3-Methyl-6-pyridin-4-yl-2-[3-(3-pyridin-4-yl-propoxy)-benzylamino]-3H-pyrimidin-4-one,
3-(6-Amino-hexyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
15 3-(6-Amino-hexyl)-2-phenethylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-(2-Amino-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-(3-Amino-propyl)-2-[2-(2-methoxy-phenyl)-ethylamino]-6-pyridin-4-yl-3H-
pyrimidin-4-one,
3-(2-Hydroxy-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
20 3-(2-Dimethylamino-ethyl)-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-
one,
3-(5-Amino-pentyl)-2-phenethylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-(4-Amino-butyl)-2-phenethylamino-6-pyridin-4-yl-3H-pyrimidin-4-one,
25 2-(4-Amino-butylamino)-3-benzyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
2-(6-Amino-hexylamino)-3-benzyl-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(2-phenylthio-ethylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(2-phenylamino-ethylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(2-phenoxy-ethylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
3-Methyl-2-(3-phenyl-allylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one,
30 3-Methyl-2-(3-phenyl-propylamino)-6-pyridin-4-yl-3H-pyrimidin-4-one, and
3-Methyl-6-pyridin-4-yl-2(2-pyridin-2-yl-ethylamino)-3H-pyrimidin-4-one
or a salt thereof, or a solvate thereof or a hydrate thereof.

9. A medicament comprising as an active ingredient a substance selected
35 from the group consisting of a pyrimidone derivative according to claim 1.

10. A GSK3 β inhibitor selected from the group of a pyrimidone derivative
represented by formula (I) or salts thereof, or a solvate thereof or a hydrate thereof

according to claim 1.

11. Use of a compound according to claim 1 for the preparation of a medicament for preventive and/or therapeutic treatment of a disease caused by
5 abnormal GSK3 β activity.

12. Use of a compound according to claims 1 to 8 for the preparation of a medicament for preventive and/or therapeutic treatment of a neurodegenerative disease.

10

13. Use of a compound according to claim 12, wherein the disease is selected from the group consisting of Alzheimer's disease, Parkinson's disease, tauopathies, vascular dementia; acute stroke, traumatic injuries; cerebrovascular accidents, brain cord trauma, spinal cord trauma; peripheral neuropathies; 15 retinopathies or glaucoma.

14. Use of a compound according to claims 1 to 8 for the preparation of a medicament for preventive and/or therapeutic treatment of non-insulin dependent diabetes; obesity; manic depressive illness; schizophrenia; alopecia; or cancers.

20

15. Use according to claim 14 wherein cancer is breast cancer, non-small cell lung carcinoma, thyroid cancer, T or B-cell leukemia or virus-induced tumors.

INTERNATIONAL SEARCH REPORT

Intern. Application No
PCT/EP 01/03640

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D401/04 A61K31/505 A61P25/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 00 18758 A (MITSUBISHI CHEMICAL CORP.) 6 April 2000 (2000-04-06) claims 1-12 —	1-15
A	CHEMICAL ABSTRACTS, vol. 84, no. 7, 16 February 1976 (1976-02-16) Columbus, Ohio, US; abstract no. 44112b, page 502; XP002127060 abstract & JP 07 435631 A (KOWA CO., LTD.) 25 September 1974 (1974-09-25) — —/—	1-9

Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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- *E* earlier document but published on or after the international filing date
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- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

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Date of the actual completion of the international search

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	CHEMICAL ABSTRACTS, vol. 83, no. 1, 7 July 1975 (1975-07-07) Columbus, Ohio, US; abstract no. 10127z, page 853; XP002127062 abstract & JP 07 435633 A (H. MORI) 25 September 1974 (1974-09-25) ---	1-9
A	WO 98 24780 A (AMGEN INC.) 11 June 1998 (1998-06-11) claims 1-31 ---	1-15
A	US 4 460 589 A (W. WIERENGA, H. I. SKULNICK) 17 July 1984 (1984-07-17) claims 1-4 ---	1-15
A	H. I. SKULNICK ET AL.: "Pyrimidinones. 3. N-Substituted 6-Phenylpyrimidinones and Pyrimidinediones with Diuretic/Hypotensive and Antiinflammatory Activity" J. MED. CHEM., vol. 29, no. 8, 1986, pages 1499-1504, XP000919430 table I ---	1-15
A	H. I. SKULNICK ET AL.: "Pyrimidinones. 1. 2-Amino-5-halo-6-aryl-4(3H)-pyrimidinones. Interferon-Inducing Antiviral Agents" J. MED. CHEM., vol. 28, no. 12, 1985, pages 1864-1869, XP002127057 table I ---	1-15
A	H.-J. KABBE: "Substituierte 4-Hydroxy- und 4-Amino-pyrimidine" LIEBIGS ANN. CHEM., vol. 704, 1967, pages 144-149, XP002127058 table I ---	1-15

INTERNATIONAL SEARCH REPORT

Information on patent family members

Interr. Serial Application No.

PCT/EP 01/03640

Patent document cited in search report		Publication date		Patent family member(s)	Publication date
WO 0018758	A	06-04-2000	AU	5759999 A	17-04-2000
JP 7435631	A			NONE	
JP 7435633	A			NONE	
WO 9824780	A	11-06-1998	AU	5525498 A	29-06-1998
			AU	733877 B	31-05-2001
			AU	6012098 A	29-06-1998
			BG	103512 A	31-07-2000
			BG	103521 A	31-07-2000
			BR	9713850 A	29-02-2000
			BR	9713863 A	14-03-2000
			CN	1246857 A	08-03-2000
			CN	1246858 A	08-03-2000
			CZ	9902015 A	17-11-1999
			CZ	9902016 A	17-11-1999
			EP	0948496 A	13-10-1999
			EP	0948497 A	13-10-1999
			HU	0001140 A	28-04-2001
			HU	0001698 A	28-04-2001
			WO	9824782 A	11-06-1998
			US	6096753 A	01-08-2000
US 4460589	A	17-07-1984	US	4593096 A	03-06-1986

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